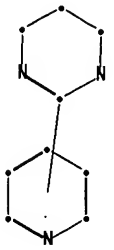
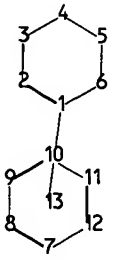
C:\STNEXP4\QUERIES\10009477 (species).str




ring nodes :
1 2 3 4 5 6 7 8 9 10 11 12
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12
exact/norm bonds :
1-2 1-6
exact bonds :
2-3 3-4 4-5 5-6
normalized bonds :
7-8 7-12 8-9 9-10 10-11 11-12
isolated ring systems :
containing 1 : 7 :

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom
12:Atom 13:CLASS

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=>
Uploading 10009477 (species).str

L1      STRUCTURE UPLOADED

=> d l1
L1 HAS NO ANSWERS
L1              STR
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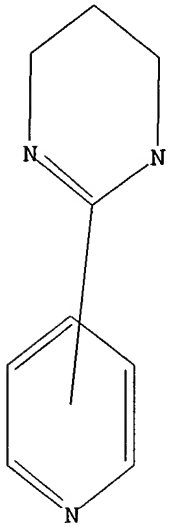


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Structure attributes must be viewed using STN Express query preparation.


=> s l1 sss sam
SAMPLE SEARCH INITIATED 11:39:59 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED -  19526 TO ITERATE

  5.1% PROCESSED    1000 ITERATIONS                                10 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:       382174 TO   398866
PROJECTED ANSWERS:            3067 TO     4743

L2           10 SEA SSS SAM L1

=>
Uploading 10009477 (species).str

L3      STRUCTURE UPLOADED

=> d l3
L3 HAS NO ANSWERS
L3              STR
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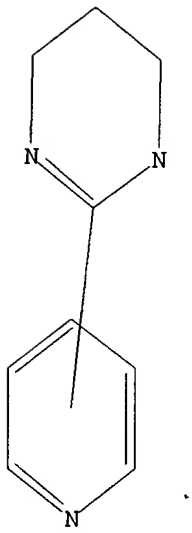
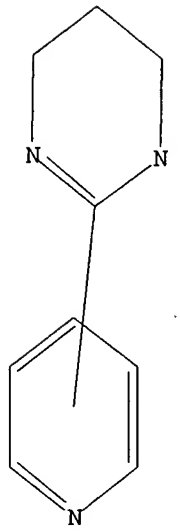



Structure attributes must be viewed using STN Express query preparation.

```
=> s l3 sss sam
SAMPLE SEARCH INITIATED 11:41:11 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED -   8547 TO ITERATE

 11.7% PROCESSED    1000 ITERATIONS                                0 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:       165404 TO   176476
PROJECTED ANSWERS:               0 TO        0

L4            0 SEA SSS SAM L3

=>
Uploading 10009477 (species).str

L5      STRUCTURE UPLOADED

=> d l5
L5 HAS NO ANSWERS
L5              STR
```




Structure attributes must be viewed using STN Express query preparation.

```
=> s l5 sss sam
SAMPLE SEARCH INITIATED 11:42:18 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED -  19526 TO ITERATE

  5.1% PROCESSED    1000 ITERATIONS                                0 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:       382174 TO   398866
PROJECTED ANSWERS:               0 TO        0

L6            0 SEA SSS SAM L5

=> s l5 sss ful
FULL SEARCH INITIATED 11:42:30 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 396496 TO ITERATE

100.0% PROCESSED  396496 ITERATIONS                             49 ANSWERS
SEARCH TIME: 00.00.05

L7           49 SEA SSS FUL L5

=> s l7
L8          20 L7

=> d l8 1-20 bib,ab,hitstr
```

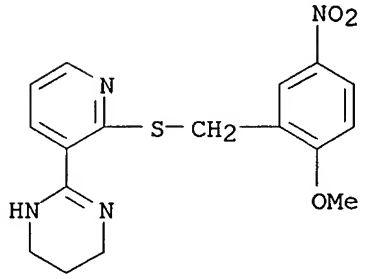
L8 ANSWER 1 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 2002:615577 CAPLUS
DN 137:169536
TI Preparation of aryl-substituted tetrahydropyrimidines and related compounds as melanocortin-4 receptor binding compounds
IN Maguire, Martin P.; Dai, Mingshi; Vos, Tricia J.
PA Millennium Pharmaceuticals, Inc., USA
SO PCT Int. Appl., 228 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|
| PI WO 2002062766 | A2 | 20020815 | WO 2002-US3566 | 20020207 |

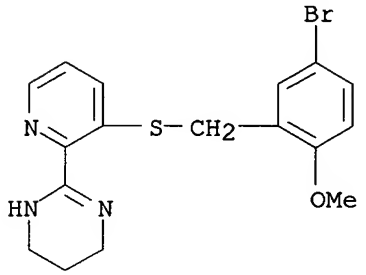
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRAI US 2001-778468 A 20010207 ← I have this case.
OS MARPAT 137:169536
AB Title compds. I [wherein A and B = independently (un)substituted biaryl, (hetero)aryl, Ph, (cyclo)alkyl, (cyclo)alkoxy, alkenyl, alkynyl, OH, acyl(oxy), carbamoyl, amino, thiol, amidino, imino, NO2, N3, etc.; L1 and L2 =- covalent bond or (un)substituted alkyl optionally interrupted by O, S, or N; r = covalent bond, CH, CH2, CHR1, CR1R2, or H; t = CH, CH2, CHR3, CR3R4, or H; s = CHR5, CR5R6, or absent; R = H, (un)substituted alkyl, arylalkyl, or heteroalkyl, and may optionally be linked to A, B, L1, or L2; R1-R6 = independently (un)substituted alkyl, halo, thiol, thioether, thioalkyl, alkoxy, and may be optionally linked to each other to form addnl. ring moieties, e.g., quinoxalinyl; or pharmaceutically acceptable salts thereof] were prepd. as melanocortin-4 receptor binding (MC4-R) compds. For example, stirring a soln. of .alpha.-tolunitrile with diisopropylamine and BuLi in hexanes at -78.degree. under nitrogen for 1 h, followed by addn. of HMPA and 1-chloromethylnaphthalene in THF, afforded 2-(2-naphthalen-1-ylethyl)benzonitrile. Heating the benzonitrile with 1,3-diaminopropane in the presence of H2S at 80.degree. for 72 h gave the tetrahydropyrimidinyl cycloaddn. product II. The latter exhibited exemplary inhibition of MC4-R in a scintillation proximity assay. I are useful for the treatment of disorders assocd. with pigmentation, bones, or wt. loss (no data).
IT **325800-74-6P**, 2-[2-(2-Methoxy-5-nitrobenzylsulfanyl)pyridin-3-yl]-1,4,5,6-tetrahydropyrimidine **325823-83-4P**, 2-[3-(5-Bromo-2-methoxybenzylsulfanyl)pyridin-2-yl]-1,4,5,6-tetrahydropyrimidine **326483-15-2P 447465-95-4P**
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(MC4-R binding compd.; prepn. of aryl-substituted tetrahydropyrimidines and related compds. as melanocortin-4 receptor binding compds. for treatment of pigmentation, bone, and wt. loss disorders)
RN 325800-74-6 CAPLUS

CN Pyrimidine, 1,4,5,6-tetrahydro-2-[2-[[(2-methoxy-5-nitrophenyl)methyl]thio]-3-pyridinyl]- (9CI) (CA INDEX NAME)



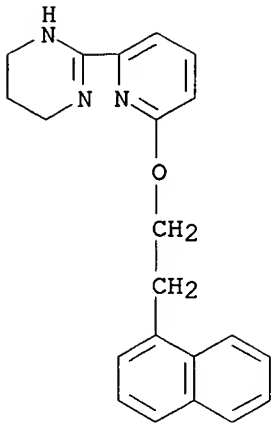

RN 325823-83-4 CAPLUS
CN Pyrimidine, 2-[3-[[(5-bromo-2-methoxyphenyl)methyl]thio]-2-pyridinyl]-1,4,5,6-tetrahydro- (9CI) (CA INDEX NAME)




RN 326483-15-2 CAPLUS
CN Formic acid, compd. with 1,4,5,6-tetrahydro-2-[6-[2-(1-naphthalenyl)ethoxy]-2-pyridinyl]pyrimidine (1:1) (9CI) (CA INDEX NAME)

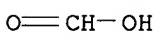
CM 1

CRN 326483-14-1
CMF C21 H21 N3 O

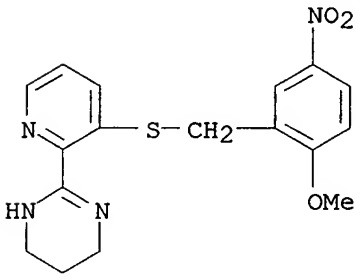


CM 2

CRN 64-18-6
CMF C H2 O2



RN 447465-95-4 CAPLUS
CN Pyrimidine, 1,4,5,6-tetrahydro-2-[3-[[(2-methoxy-5-nitrophenyl)methyl]thio]-2-pyridinyl]-, monohydrobromide (9CI) (CA INDEX NAME)



● HBr

L8 ANSWER 2 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 2001:793434 CAPLUS
DN 135:339275
TI Cyclic amidines, nicotinic acetylcholine .alpha.4.beta.2 receptor activators containing them, and pharmaceuticals
IN Imoto, Masahiro; Iwanami, Tatsuya; Akabane, Minako; Tani, Yoshihiro
PA Suntory, Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 25 pp.
CODEN: JKXXAF
DT Patent
LA Japanese
FAN.CNT 1

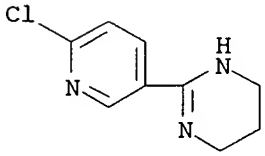
| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|---|
| PI | JP 2001302643 | A2 | 20011031 | JP 2000-120976 | 20000421 |
| | WO 2001081334 | A2 | 20011101 | WO 2001-JP3378 | 20010420 |
| | WO 2001081334 | A3 | 20020808 | | |

W: AU, CA, CN, KR, US
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR

PRAI JP 2000-120976 A 20000421
OS MARPAT 135:339275
AB The activators, useful for treatment of brain function disorders, contain cyclic amidines I [A1, A2 = H, (un)substituted alkyl, (un)substituted aryl, (un)substituted heterocyclyl; X = (un)substituted C2H4, (un)substituted CH:CH, (un)substituted (CH2)3, (un)substituted CH2CH2NH] or their salts. Trimethylenediamine was cyclocondensed with Et (6-chloro-3-pyridyl)acetate and treated with fumaric acid to give I · fumarate (A1 = H, A2 = 6-chloro-3-pyridylmethyl, X = CH:CH), which showed affinity with rat nicotinic acetylcholine .alpha.4.beta.2 receptor with Ki of 29 nM, vs. 1.6 nM, for nicotine. Pharmaceutical formulations contg. I are given.
IT **371121-78-7P 371121-82-3P**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of cyclic amidines as nicotinic acetylcholine .alpha.4.beta.2 receptor activators)
RN 371121-78-7 CAPLUS
CN Pyrimidine, 2-(6-chloro-3-pyridinyl)-1,4,5,6-tetrahydro-, (2E)-2-butenedioate (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 371121-77-6
CMF C9 H10 Cl N3

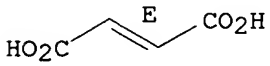



Elected species.

CM 2

CRN 110-17-8
CMF C4 H4 O4

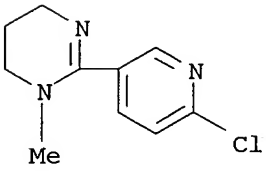
Double bond geometry as shown.



RN 371121-82-3 CAPLUS
CN Pyrimidine, 2-(6-chloro-3-pyridinyl)-1,4,5,6-tetrahydro-1-methyl-, ethanedioate (1:1) (9CI) (CA INDEX NAME)

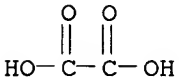
CM 1

CRN 371121-81-2
CMF C10 H12 Cl N3

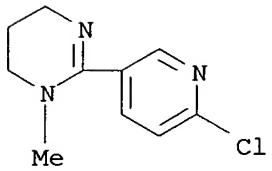


CM 2

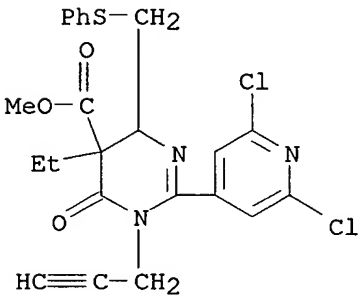
CRN 144-62-7
CMF C2 H2 O4



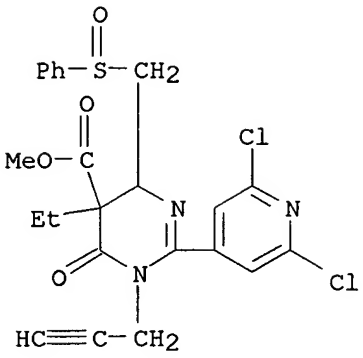
IT **371121-81-2**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(prepn. of cyclic amidines as nicotinic acetylcholine .alpha.4.beta.2 receptor activators)
RN 371121-81-2 CAPLUS
CN Pyrimidine, 2-(6-chloro-3-pyridinyl)-1,4,5,6-tetrahydro-1-methyl- (9CI) (CA INDEX NAME)



L8 ANSWER 4 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1997:417074 CAPLUS
DN 127:149117
TI Synthesis of 2-(2,6-dichloro-4-pyridyl)-3-propargyl-5-ethyl-6-methyl-4(3H)-pyrimidinone, a promising new herbicide
AU Taylor, Edward C.; Zhou, Ping; Tice, Colin M.; Lidert, Zev; Roemmele, Renee C.
CS Dep. Chemistry, Princeton Univ., Princeton, NJ, 08544, USA
SO Tetrahedron Letters (1997), 38(25), 4339-4342
CODEN: TELEAY; ISSN: 0040-4039
PB Elsevier
DT Journal
LA English
OS CASREACT 127:149117
AB A novel synthesis of the title compd. I, a promising new herbicide, is reported which features regioselective carbon, followed by nitrogen, dialkylation of an intermediate dianion, and a tandem one-pot sequence of reactions involving sigmatropic sulfoxide elimination, LiCl-induced demethylation of a carbomethoxy grouping, decarboxylation, and isomerization/aromatization.
IT **193286-89-4P 193286-90-7P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate in prepn. of (dichloropyridyl)propargylethylmethylpyrimidinone)
RN 193286-89-4 CAPLUS
CN 5-Pyrimidinecarboxylic acid, 2-(2,6-dichloro-4-pyridinyl)-5-ethyl-1,4,5,6-tetrahydro-6-oxo-4-[(phenylthio)methyl]-1-(2-propynyl)-, methyl ester (9CI) (CA INDEX NAME)



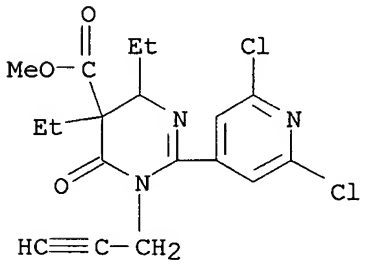

RN 193286-90-7 CAPLUS
CN 5-Pyrimidinecarboxylic acid, 2-(2,6-dichloro-4-pyridinyl)-5-ethyl-1,4,5,6-tetrahydro-6-oxo-4-[(phenylsulfinyl)methyl]-1-(2-propynyl)-, methyl ester (9CI) (CA INDEX NAME)



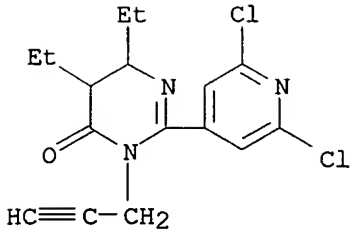
L8 ANSWER 5 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1996:630487 CAPLUS
DN 125:275903
TI Preparation of 2-aryl-5,6-dihydropyrimidin-4-ones as herbicides
IN Tice, Colin Michael; Bryman, Lois Merle; Roemmele, Renee Caroline
PA Rohm and Haas Company, USA
SO Eur. Pat. Appl., 24 pp.
CODEN: EPXXDW
DT Patent
LA English
FAN.CNT 1

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|
| PI EP 733622 | A1 | 19960925 | EP 1996-301548 | 19960306 |
| R: DE, ES, FR, GB, IT | | | | |
| US 5629264 | A | 19970513 | US 1995-409293 | 19950323 |
| AU 9648005 | A1 | 19961003 | AU 1996-48005 | 19960312 |
| AU 710045 | B2 | 19990909 | | |
| CA 2171926 | AA | 19960924 | CA 1996-2171926 | 19960315 |
| BR 9601059 | A | 19980106 | BR 1996-1059 | 19960320 |
| CN 1141292 | A | 19970129 | CN 1996-103157 | 19960321 |
| JP 08283245 | A2 | 19961029 | JP 1996-91856 | 19960322 |
| PRAI US 1995-409293 | | 19950323 | | |

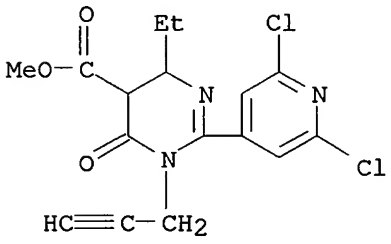
OS MARPAT 125:275903
AB Title compds. [I; R = (hetero)aryl; R3 = (halo)alkenyl, alkoxyalkyl,cyanoalkyl, etc.; R5a,R5b = H, halo, alkyl, alkoxy, etc.; R6a,R6b = H, halo, alkyl, alkoxy, etc.; R5aR6a = (CH2)2-5; X = O or S] were prepd. Thus, 3-ClC6H4C(:NH)NH2 was cyclocondensed with F3CCH:CHCO2Et and the product N-alkylated with HC.tplbond.CCH2Br to give I (R = C6H4Cl-3, R3 = CH2C.tplbond.CH, R5a = R5b = R6b = H, R6a = CF3) which gave 90-100% control of 5 weeds at 1200g/ha preemergent.
IT **182254-57-5P 182254-58-6P 182254-59-7P 182254-62-2P**
RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of 2-aryl-5,6-dihydropyrimidin-4-ones as herbicides)
RN 182254-57-5 CAPLUS
CN 5-Pyrimidinecarboxylic acid, 2-(2,6-dichloro-4-pyridinyl)-4,5-diethyl-1,4,5,6-tetrahydro-6-oxo-1-(2-propynyl)-, methyl ester (9CI) (CA INDEX NAME)



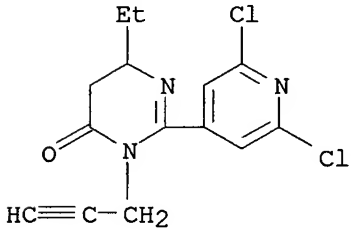

RN 182254-58-6 CAPLUS
CN 4(3H)-Pyrimidinone, 2-(2,6-dichloro-4-pyridinyl)-5,6-diethyl-5,6-dihydro-3-(2-propynyl)- (9CI) (CA INDEX NAME)



RN 182254-59-7 CAPLUS
CN 5-Pyrimidinecarboxylic acid, 2-(2,6-dichloro-4-pyridinyl)-4-ethyl-1,4,5,6-tetrahydro-6-oxo-1-(2-propynyl)-, methyl ester (9CI) (CA INDEX NAME)



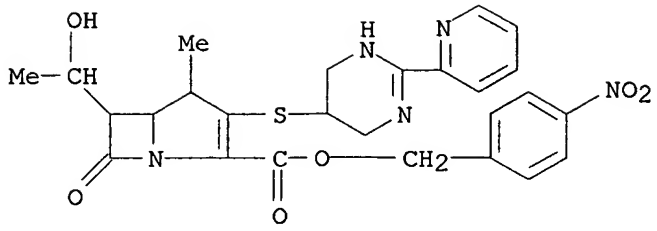
RN 182254-62-2 CAPLUS
CN 4(3H)-Pyrimidinone, 2-(2,6-dichloro-4-pyridinyl)-6-ethyl-5,6-dihydro-3-(2-propynyl)- (9CI) (CA INDEX NAME)



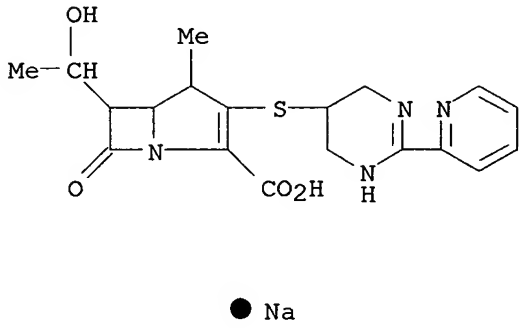
L8 ANSWER 6 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1996:625489 CAPLUS
DN 125:275528
TI Preparation of carbapenem compounds as antibacterials
IN Miwa, Tetsuo; Higuchi, Noriko; Soejima, Seizo; Okonogi, Kenji
PA Takeda Chemical Industries, Ltd., Japan
SO PCT Int. Appl., 152 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|---|
| PI | WO 9626939 | A1 | 19960906 | WO 1996-JP509 | 19960301 |
| | W: AL, AM, AU, AZ, BB, BG, BR, BY, CA, CN, CZ, EE, GE, HU, IS, KG, KR, KZ, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TR, TT, UA, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU | | | | |
| | RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG | | | | |
| | AU 9648448 | A1 | 19960918 | AU 1996-48448 | 19960301 |
| | JP 09165388 | A2 | 19970624 | JP 1996-44424 | 19960301 |
| PRAI | JP 1995-42765 | | 19950302 | | |
| | JP 1995-69343 | | 19950328 | | |
| | JP 1995-261371 | | 19951009 | | |
| | WO 1996-JP509 | | 19960301 | | |

OS MARPAT 125:275528
AB Carbapenem compds. I [R1 = (un)substituted lower alkyl group; R2 = H, lower alkyl group; Y = bond, (un)substituted alkylene group; W = Q, Q1; A = (un)hydrogenated pyrimidine ring which may be substituted; X = (un)substituted hetero-atom; X may form a ring, taken together with the ring A-constituent nitrogen atom; X1 = O, S; X2 = NH, O or S; ring A1 may be substituted, provided that, when X2 = NH, X1 = O; R2 = lower alkyl] or their esters or salts, useful as antibacterials, are prepd. Thus, 5-(4-methoxybenzylthio)-2-(methylthio)-1,4,5,6-tetrahydropyrimidine was treated with CF3COOH-anisole and the resulting pyrimidinethiol was reacted with 4-nitrobenzyl (4R,5R,6S)-3-[(diphenylphosphono)oxy]-6-[(R)-1-hydroxyethyl]-4-methyl-7-oxo-1-azabicyclo[3.2.0]hept-2-ene-2-carboxylate in MeCN contg. diisopropylethylamine to give the title compd. 4-nitrobenzyl (4R,5R,6S)-3-[2-(methylthio)-1,4,5,6-tetrahydropyrimidin-5-ylthio]-6-[(R)-1-hydroxyethyl]-4-methyl-7-oxo-1-azabicyclo[3.2.0]hept-2-ene-2-carboxylate, which was sapond. with NaOH to give sodium (4R,5S,6S)-[(R)-1-hydroxyethyl]-4-methyl-3-[(2-methylthio-1,4,5,6-tetrahydropyrimidin-5-yl)thio]-7-oxo-1-azabicyclo[3.2.0]hept-2-ene-2-carboxylate. In an in vitro study using Muller-Hinton agar medium, this had an IC50 of 0.05.times.106 CFU/mL against Escherichia coli vs. 0.1.times.106 CFU/mL for Impenem.
IT **182203-13-0P**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(prepn. of carbapenem compds. as antibacterials)
RN 182203-13-0 CAPLUS
CN 1-Azabicyclo[3.2.0]hept-2-ene-2-carboxylic acid, 6-(1-hydroxyethyl)-4-methyl-7-oxo-3-[[1,4,5,6-tetrahydro-2-(2-pyridinyl)-5-pyrimidinyl]thio]-, (4-nitrophenyl)methyl ester (9CI) (CA INDEX NAME)

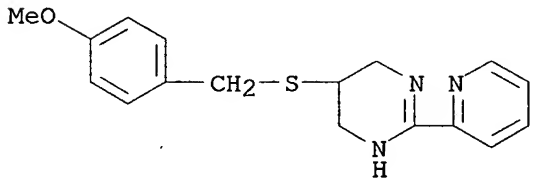


IT **182203-15-2P**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of carbapenem compds. as antibacterials)
RN 182203-15-2 CAPLUS
CN 1-Azabicyclo[3.2.0]hept-2-ene-2-carboxylic acid, 6-(1-hydroxyethyl)-4-methyl-7-oxo-3-[[1,4,5,6-tetrahydro-2-(2-pyridinyl)-5-pyrimidinyl]thio]-, monosodium salt (9CI) (CA INDEX NAME)



● Na

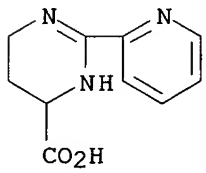
IT **182204-23-5P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of carbapenem compds. as antibacterials)
RN 182204-23-5 CAPLUS
CN Pyrimidine, 1,4,5,6-tetrahydro-5-[[(4-methoxyphenyl)methyl]thio]-2-(2-pyridinyl)- (9CI) (CA INDEX NAME)



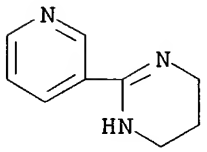
L8 ANSWER 7 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1991:632273 CAPLUS
DN 115:232273
TI Nitrogen-containing heterocyclic compounds and their optically active isomers
IN Matsumura, Koichi; Mano, Mitsuhiko; Nishimura, Tatsuo; Sugyama, Yoshio
PA Takeda Chemical Industries, Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 9 pp.
CODEN: JKXXAF
DT Patent
LA Japanese
FAN.CNT 1

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|
| PI JP 03086867 | A2 | 19910411 | JP 1990-168795 | 19900626 |
| PRAI JP 1989-163482 | | 19890626 | | |

AB Title compds. I or II (R1 = H, C1-18 alkyl, aryl, aralkyl, heterocyclic group; R2 = H, C1-8 alkyl; n = 0-3; R2 = C1-8 alkyl when n = 1 and R1 = H or Me) and their salts and optically active isomers, useful as intermediates for pharmaceuticals, agrochems., and liq. crystals, are prepd. Thus, refluxing Me propionimidate hydrochloride and 1,4-diaminobutyric acid in MeOH in the presence of NaOMe gave 90% 2-ethyl-1,4,5,6-tetrahydro-4-pyrimidinecarboxylic acid.
IT **137023-65-5P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of, by condensation of diaminobutyric acid and imidate ester)
RN 137023-65-5 CAPLUS
CN 4-Pyrimidinecarboxylic acid, 1,4,5,6-tetrahydro-2-(2-pyridinyl)- (9CI) (CA INDEX NAME)

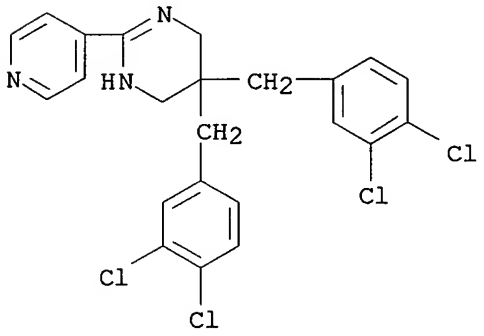


L8 ANSWER 9 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1988:32483 CAPLUS
DN 108:32483
TI Effects of nucleus basalis lesions on the muscarinic and nicotinic modulation of [3H]acetylcholine release in the rat cerebral cortex
AU Meyer, Edwin M.; Arendash, Gary W.; Judkins, Jennifer H.; Ying, Lily; Wade, Cathy; Kem, William R.
CS Sch. Med., Univ. Florida, Gainesville, FL, 32610, USA
SO J. Neurochem. (1987), 49(6), 1758-62
CODEN: JONRA9; ISSN: 0022-3042
DT Journal
LA English
AB Adult male rats received unilateral infusions of ibotenic acid (5 .mu.g/.mu.L) in the nucleus basalis magnocellularis (nbm). Two weeks later, cerebral cortical cholinergic markers (choline acetyltransferase activity, high-affinity choline uptake, and coupled acetylcholine (Ach) synthesis) were reduced in synaptosomes prepd. from the lesioned hemispheres compared to contralateral controls. The depolarization-induced release of [3H]ACh from these synaptosomes was also reduced in the lesioned hemispheres, reflecting the reduced synthesis of transmitter. However, the nbm lesions had no effect on the inhibition of release induced by 100 .mu.M oxotremorine. Synaptosomal [3H]ACh release was not altered by nicotine or the nicotinic agonists anabaseine and 2-(3-pyridyl)-1,4,5,6-tetrahydropyrimidine. Nicotine (10-100 .mu.M) did increase [3H]ACh release in control and lesioned hemispheres in cortical minces, but to a similar extent. Apparently, neither muscarinic nor nicotinic receptors modulating ACh release reside on nbm-cholinergic terminals.
IT **112147-36-1**
RL: BIOL (Biological study)
(acetylcholine release response to, in brain cerebral cortex, nucleus basalis in relation to)
RN 112147-36-1 CAPLUS
CN Pyrimidine, 1,4,5,6-tetrahydro-2-(3-pyridinyl)- (9CI) (CA INDEX NAME)




same as #13

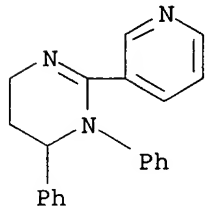
L8 ANSWER 10 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1983:463892 CAPLUS
DN 99:63892
TI Appraisals of compounds of diverse chemical classes for capacities to cure infections with sporozoites of Plasmodium cynomolgi
AU Schmidt, L. H.
CS South. Res. Inst., Kettering-Meyer Lab., Birmingham, AL, USA
SO Am. J. Trop. Med. Hyg. (1983), 32(2), 231-57
CODEN: AJTHAB; ISSN: 0002-9637
DT Journal
LA English
AB Compds. of widely diverse structures were appraised for radical curative activity in rhesus monkeys infected with sporozoites of the B strain of P. cynomolgi, using an evaluation system that provided a preliminary assessment with from 0.1-1.0 g of compd. and tests against 1 to 5 active infections. None of 32 compds. in a misc. structure category, none of 7 agents of antibiotic origin, none of twelve 1,5-naphthyridines, and none of seven 7-aminoquinolines exhibited curative activity at the largest test doses. One of 12 newly synthesized pyrocatechols appeared to have curative effect. Two of twenty 6-aminoquinolines showed curative effect at or near max. tolerated doses. In contrast, 90 of 174 8-aminoquinolines had curative effects, 18 of the 90 being as active as primaquine, 8 twice as active, and 6 four times as active. There were major disaggreements between the above results and those recorded by others in mice inoculated with sporozites of P. berghei or P. yoelii nigeriensis. These discrepancies were of serious dimensions in evaluations of the 8-aminoquinolines. This, plus previous near flawless performances of P. cynomolgi in identifying agents that would cure naturally acquired P. vivax infections, led to the suggestion that the abbreviated simian model employed in these studies be used hereafter in primary screening of new agents for radical curative activity.
IT **80061-38-7**
RL: BIOL (Biological study)
(Plasmodium cynomolgi sporozoites infection therapy with)
RN 80061-38-7 CAPLUS
CN Pyrimidine, 5,5-bis[(3,4-dichlorophenyl)methyl]-1,4,5,6-tetrahydro-2-(4-pyridinyl)- (9CI) (CA INDEX NAME)



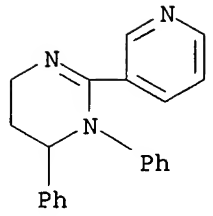
L8 ANSWER 11 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1983:422491 CAPLUS
DN 99:22491
TI 1,4,5,6-Tetrahydropyrimidine derivatives
IN Gauthier, Jean A.; Jirkovsky, Ivo
PA Ayerst, McKenna and Harrison Ltd., Can.
SO U.S., 12 pp.
CODEN: USXXAM
DT Patent
LA English
FAN.CNT 1

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|
| US 4379926 | A | 19830412 | US 1978-904124 | 19780508 |

OS CASREACT 99:22491
AB Diphenylpyrimidines I (R = alkyl, Ph, 2-furyl, 3-pyridinyl, 2-thienyl, dialkylamino, ZR2; R2 = 1-piperidinyl, 4-morpholinyl; Z = alkylene) were prepd. by cyclization of PhNHCHPhCH2CH2COR (II). Thus, PhNHNH2 108.14, CH2O 30.03, and styrene 104.1 g were cyclocondensed to give 135.9 g 2,3-diphenylpyrazolidine. The latter compd. (133.9 g) was hydrogenated to give 48.4 g PhNHCHPhCH2CH2NH2, which (5.0 g) was benzoylated to give 2.0 g II (R = Ph). This (12.0 g) was cyclized with POCl3 to give 9.3 g I (R = Ph) (III). In rats, 6.25 mg III/kg orally was an effective diuretic.
IT **86203-86-3P**
RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. and diuretic activity of)
RN 86203-86-3 CAPLUS
CN Pyrimidine, 1,4,5,6-tetrahydro-1,6-diphenyl-2-(3-pyridinyl)- (9CI) (CA INDEX NAME)

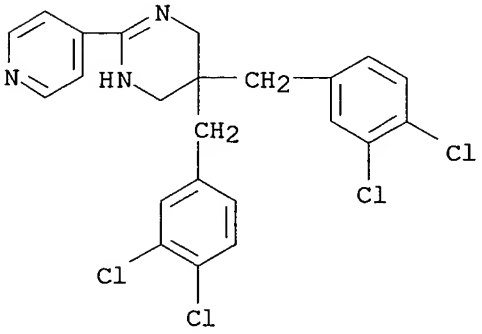



IT **86203-87-4P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)
RN 86203-87-4 CAPLUS
CN Pyrimidine, 1,4,5,6-tetrahydro-1,6-diphenyl-2-(3-pyridinyl)-, monohydrobromide (9CI) (CA INDEX NAME)




● HBr

L8 ANSWER 13 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1982:62629 CAPLUS
DN 96:62629
TI New tissue schizontocidal antimalarial drugs
AU Davidson, David E., Jr.; Ager, Arba L.; Brown, John L.; Chapple, Frank E.; Whitmire, Richard E.; Rossan, Richard N.
CS Dep. Parasitol., Walter Reed Army Inst. Res., Washington, DC, 20012, USA
SO Bull. W. H. O. (1981), 59(3), 463-79
CODEN: BWHOA6; ISSN: 0366-4996
DT Journal
LA English
AB Over 700 causal prophylactic and radical curative antimalarial drugs have been discovered during the screening of approx. 4000 chem. compds. in rodent and simian malaria models. Causal prophylactic activity in the Plasmodium berghei-rodent model was demonstrated by 10 distinct groups of chems.: 1) tetrahydrofolate dehydrogenase inhibitors, 2) naphthoquinones, 3) dihydroacridinediones, 4) tetrahydrofurans, 5) guanylhydrazones, 6) clopidol analogs, 7) quinoline esters, 8) dibenzyltetrahydropyrimidines, 9) 6-aminoquinolines, 10) 8-aminoquinolines. Of the causal prophylactic compds., only the 6- and 8-aminoquinolines were capable of curing persistent exoerythrocytic infections of P. cynomolgi in rhesus monkeys. The 6-aminoquinolines were substantially less active than primaquine. A series of 4-methyl-5-phenoxy-6-methoxy-8-aminoquinolines I (R = F, Cl, OMe, CF3) were potent blood schizontocides and radical curative drugs. The most active member of this series WR 225448 (I succinate salt, R = CF3) [80065-56-1], was 5 times more active than primaquine in curing persistent exoerythrocytic infections of P. cynomolgi in rhesus monkeys. As a blood schizontocide, WR 225448 was effective in animal models against P. berghei, P. cynomolgi, P. vivax, and both drug-sensitive and drug-resistant strains of P. falciparum. WR 225448 was also more toxic than primaquine in rats during subacute (28-day) administration.
IT **80061-38-7**
RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(antimalarial activity of)
RN 80061-38-7 CAPLUS
CN Pyrimidine, 5,5-bis[(3,4-dichlorophenyl)methyl]-1,4,5,6-tetrahydro-2-(4-pyridinyl)- (9CI) (CA INDEX NAME)




Same as #10

L8 ANSWER 14 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1981:103366 CAPLUS
DN 94:103366
TI Urea and amido compounds
IN Marxer, Adrian
PA Ciba-Geigy A.-G., Switz.
SO S. African, 34 pp.
CODEN: SFXXAB
DT Patent
LA English
FAN.CNT 2

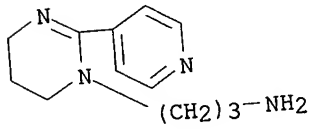
| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|---|
| PI | ZA 7901062 | A | 19800326 | ZA 1979-1062 | 19790307 |
| | CA 1125759 | A1 | 19820615 | CA 1979-321545 | 19790215 |
| | US 4292429 | A | 19810929 | US 1979-14661 | 19790223 |
| | FI 7900740 | A | 19790909 | FI 1979-740 | 19790305 |
| | FI 70708 | B | 19860626 | | |
| | FI 70708 | C | 19861006 | | |
| | EP 4561 | A2 | 19791017 | EP 1979-100647 | 19790305 |
| | EP 4561 | B1 | 19811104 | | |
| | EP 4561 | A3 | 19791114 | | |
| | R: BE, CH, DE, FR, GB, IT, LU, NL, SE | | | | |
| | CS 244656 | B2 | 19860814 | CS 1979-1460 | 19790305 |
| | ES 478342 | A1 | 19790516 | ES 1979-478342 | 19790306 |
| | DD 142336 | C | 19800618 | DD 1979-211405 | 19790306 |
| | PL 116762 | B1 | 19810630 | PL 1979-213924 | 19790306 |
| | PL 123150 | B1 | 19820930 | PL 1979-221681 | 19790306 |
| | IL 56797 | A1 | 19820930 | IL 1979-56797 | 19790306 |
| | DK 7900952 | A | 19790909 | DK 1979-952 | 19790307 |
| | NO 7900765 | A | 19790911 | NO 1979-765 | 19790307 |
| | NO 152606 | B | 19850715 | | |
| | NO 152606 | C | 19851023 | | |
| | AU 7944900 | A1 | 19790913 | AU 1979-44900 | 19790307 |
| | AU 531006 | B2 | 19830804 | | |
| | AT 7901710 | A | 19810315 | AT 1979-1710 | 19790307 |
| | AT 364375 | B | 19811012 | | |
| | SU 845779 | A3 | 19810707 | SU 1979-2733999 | 19790307 |
| | HU 25271 | O | 19830628 | HU 1979-CI1920 | 19790307 |
| | HU 182940 | B | 19840328 | | |
| | JP 54125668 | A2 | 19790929 | JP 1979-26245 | 19790308 |
| | JP 62009109 | B4 | 19870226 | | |
| | SU 923367 | A3 | 19820423 | SU 1980-2872253 | 19800118 |
| | AT 8003951 | A | 19810515 | AT 1980-3951 | 19800730 |
| | AT 365179 | B | 19811228 | | |
| | US 4420619 | A | 19831213 | US 1981-247427 | 19810325 |
| | CS 244700 | B2 | 19860814 | CS 1984-8407 | 19841105 |
| PRAI | CH 1978-2519 | | 19780308 | | |
| | US 1979-14661 | | 19790223 | | |
| | CS 1979-1460 | | 19790305 | | |
| | AT 1979-1710 | | 19790307 | | |

AB The antitumor (no data) compds. I (R = aryl, arylamino, aralkyl, arylaminoalkyl; R1 = aryl, arylamino; X = O, S; X1 = alkylene n = 1, 2) were prepd. Thus, 2,6-Cl2C6H3NHCH2CN was treated with HN(CH2CH2NH2)2 to give II (R2 = H), which was treated with 4-MeC6H4NCO to give II (R2 = CONHC6H4Me-4).

IT **73998-75-1P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)

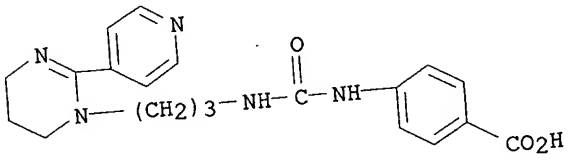
(prepn. and reaction of, with isocyanates)

RN 73998-75-1 CAPLUS
CN 1(4H)-Pyrimidinepropanamine, 5,6-dihydro-2-(4-pyridinyl)- (9CI) (CA INDEX NAME)

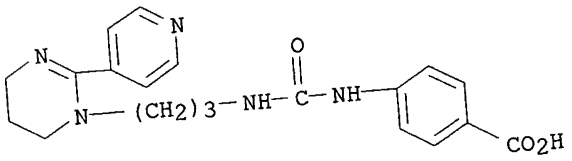


IT **73998-73-9P 76692-14-3P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 73998-73-9 CAPLUS
CN Benzoic acid, 4-[[[[3-[5,6-dihydro-2-(4-pyridinyl)-1(4H)-pyrimidinyl]propyl]amino]carbonyl]amino]- (9CI) (CA INDEX NAME)



RN 76692-14-3 CAPLUS
CN Benzoic acid, 4-[[[[3-[5,6-dihydro-2-(4-pyridinyl)-1(4H)-pyrimidinyl]propyl]amino]carbonyl]amino]-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

L8 ANSWER 15 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1980:446666 CAPLUS
DN 93:46666
TI Process for the preparation of novel imidazole urea and amido compounds
IN Marxer, Adrian
PA Ciba-Geigy A.-G., Switz.
SO Brit. UK Pat. Appl., 14 pp.
CODEN: BAXXDU
DT Patent
LA English
FAN.CNT 2

| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|---|
| PI | GB 2016011 | A | 19790919 | GB 1979-8098 | 19790307 |
| | GB 2016011 | B2 | 19820825 | | |
| | CA 1125759 | A1 | 19820615 | CA 1979-321545 | 19790215 |
| | US 4292429 | A | 19810929 | US 1979-14661 | 19790223 |
| | FI 7900740 | A | 19790909 | FI 1979-740 | 19790305 |
| | FI 70708 | B | 19860626 | | |
| | FI 70708 | C | 19861006 | | |
| | EP 4561 | A2 | 19791017 | EP 1979-100647 | 19790305 |
| | EP 4561 | B1 | 19811104 | | |
| | EP 4561 | A3 | 19791114 | | |
| | R: BE, CH, DE, FR, GB, IT, LU, NL, SE | | | | |
| | CS 244656 | B2 | 19860814 | CS 1979-1460 | 19790305 |
| | ES 478342 | A1 | 19790516 | ES 1979-478342 | 19790306 |
| | DD 142336 | C | 19800618 | DD 1979-211405 | 19790306 |
| | PL 116762 | B1 | 19810630 | PL 1979-213924 | 19790306 |
| | PL 123150 | B1 | 19820930 | PL 1979-221681 | 19790306 |
| | IL 56797 | A1 | 19820930 | IL 1979-56797 | 19790306 |
| | DK 7900952 | A | 19790909 | DK 1979-952 | 19790307 |
| | NO 7900765 | A | 19790911 | NO 1979-765 | 19790307 |
| | NO 152606 | B | 19850715 | | |
| | NO 152606 | C | 19851023 | | |
| | AU 7944900 | A1 | 19790913 | AU 1979-44900 | 19790307 |
| | AU 531006 | B2 | 19830804 | | |
| | AT 7901710 | A | 19810315 | AT 1979-1710 | 19790307 |
| | AT 364375 | B | 19811012 | | |
| | SU 845779 | A3 | 19810707 | SU 1979-2733999 | 19790307 |
| | HU 25271 | O | 19830628 | HU 1979-CI1920 | 19790307 |
| | HU 182940 | B | 19840328 | | |
| | JP 54125668 | A2 | 19790929 | JP 1979-26245 | 19790308 |
| | JP 62009109 | B4 | 19870226 | | |
| | SU 923367 | A3 | 19820423 | SU 1980-2872253 | 19800118 |
| | AT 8003951 | A | 19810515 | AT 1980-3951 | 19800730 |
| | AT 365179 | B | 19811228 | | |
| | US 4420619 | A | 19831213 | US 1981-247427 | 19810325 |
| | CS 244700 | B2 | 19860814 | CS 1984-8407 | 19841105 |
| PRAI | CH 1978-2519 | | 19780308 | | |
| | US 1979-14661 | | 19790223 | | |
| | CS 1979-1460 | | 19790305 | | |
| | AT 1979-1710 | | 19790307 | | |

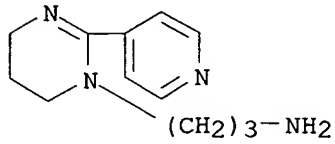
AB Ureas and amides I (R, R2 = monocyclic, carbocyclic aryl or heteroaryl; R1 = H, alkyl; n = 0, 1; m = 0, 1, 2; x = 1, 2; Z = alkylene having 2-3 C atoms in the linear chain; Z1 = O, S; Z2 = imino, bond) and I salts were prepd. E.g., 1-[2-[2-(2,6-dichloroanilinomethyl)-2-imidazolin-1-yl]ethyl]-3-(p-tolyl)urea was prepd. by stirring 1-aminoethyl-2-(2,6-dichloroanilinomethyl)-2-imidazoline with p-MeC6H4NCO in PhMe at

90.degree. for 3 h. I have a powerful action against tumors; their activities were assessed against respiratory carcinomas in golden hamsters and the Ehrlich ascites carcinoma in mice. They are particularly valuable for the treatment of bronchial carcinomas. Compns. contg. I are described.

IT **73998-75-1P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
(prepn. and addn. reaction of, with aryl isocyanate)

RN 73998-75-1 CAPLUS

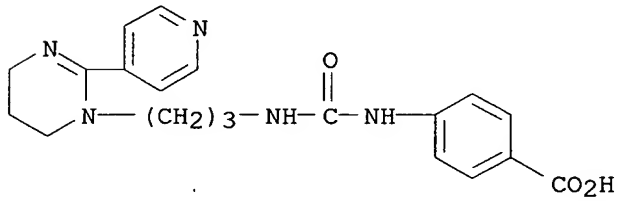
CN 1(4H)-Pyrimidinepropanamine, 5,6-dihydro-2-(4-pyridinyl)- (9CI) (CA INDEX NAME)



IT **73998-73-9P 73998-74-0P**
RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of, as neoplasm inhibitor)

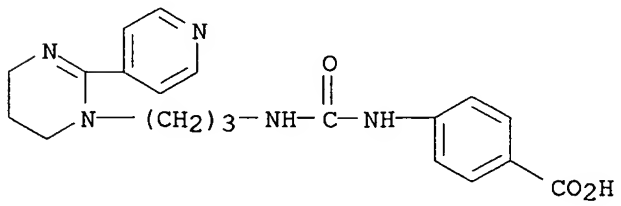
RN 73998-73-9 CAPLUS

CN Benzoic acid, 4-[[[[3-[5,6-dihydro-2-(4-pyridinyl)-1(4H)-pyrimidinyl]propyl]amino]carbonyl]amino]- (9CI) (CA INDEX NAME)



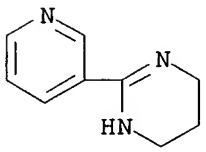
RN 73998-74-0 CAPLUS

CN Benzoic acid, 4-[[[[3-[5,6-dihydro-2-(4-pyridinyl)-1(4H)-pyrimidinyl]propyl]amino]carbonyl]amino]-, hydrochloride (9CI) (CA INDEX NAME)



● x HCl

L8 ANSWER 19 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1972:429705 CAPLUS
DN 77:29705
TI Actions of some muscarinic and nicotinic agonists on the Retzius cells of the leech
AU Woodruff, Geoffrey N.; Walker, Robert James; Newton, Lynne C.
CS Dep. Physiol. Biochem., Univ. Southampton, Southampton, Engl.
SO Comp. Gen. Pharmacol. (1971), 2(5), 106-17
CODEN: CPGPAY
DT Journal
LA English
AB Twenty-eight acetylcholine analogs, nicotinic and muscarinic agonists, and acetylenic compds. were tested on Retzius cells of the leech (Hirudo medicinalis) nerve cord. Generally the acetylcholine analogs and nicotinic agonists were powerful stimulants, causing depolarization with an increase in the rate of firing of action potentials, whereas the muscarcinic agonists usually caused hyperpolarization and inhibition of neurons when added in threshold amts. and excitation when added in relatively large amts. Carbachol [51-83-2] and 2-(3-pyridyl)-1,4,5,6-tetrahydropyrimidine-HCl (I) [**35059-05-3**] were the most potent stimulants, and [4-(m-chlorophenylcarbamoyloxy)-2-butynyl]trimethylammonium chloride (II) [55-45-8] was the most potent inhibitor.
IT **35059-05-3**
RL: PRP (Properties)
(nerve Retzius cell of leech in response to)
RN 35059-05-3 CAPLUS
CN Pyrimidine, 1,4,5,6-tetrahydro-2-(3-pyridinyl)-, monohydrochloride (9CI) (CA INDEX NAME)




● HCl

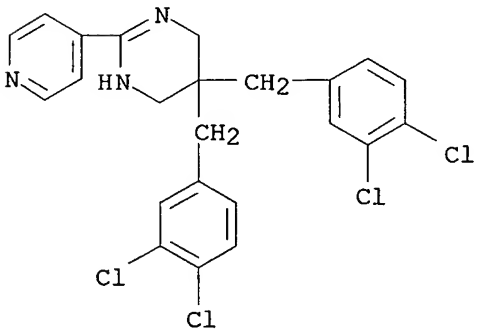
L8 ANSWER 20 OF 20 CAPLUS COPYRIGHT 2002 ACS
AN 1970:425515 CAPLUS
DN 73:25515
TI Tetrahydropyrimidines active against protozoa
PA Farbwerke Hoechst A.-G.
SO Fr. Demande, 20 pp.
CODEN: FRXXBL
DT Patent
LA French
FAN.CNT 1

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|---|---|---|---|
| PI FR 2007450 | A1 | 19700109 | FR 1969-11419 | 19690414 |
| PRAI DE 1967-1770208 | | 19680413 | | |

AB I (R = 4-pyridyl, 4-sulfamoylphenyl, 4-nitrophenyl, 3-pyridyl, 3-chloro-4nitrophenyl, 4-cyanophenyl, or a similar group; R1 = 3,4-dichlorobenzyl or 2,4-dichlorobenzyl), and their HCl, maleate, and oxalate salts, are prepd. The compds. exhibit mild activity against protozoa such as Plasmodium berghei and Babesia rodhaini. The compds. are prepd., for example, by the treatment of compds. such as 2,2-bis(3,4-dichlorobenzyl)-1,3-diaminopropane with isonicotinic acid or a similar compd. in aq. HCl.

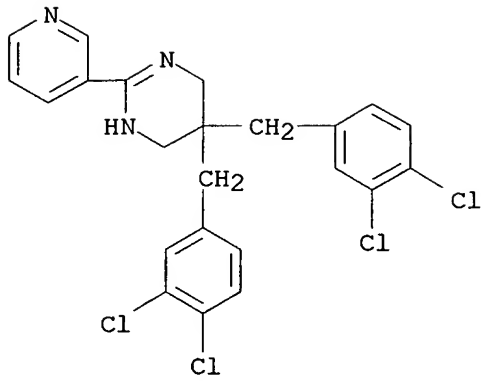
IT **27337-17-3P 27337-19-5P 27338-48-3P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 27337-17-3 CAPLUS
CN Pyrimidine, 5,5-bis(3,4-dichlorobenzyl)-1,4,5,6-tetrahydro-2-(4-pyridyl)-, dihydrochloride (8CI) (CA INDEX NAME)



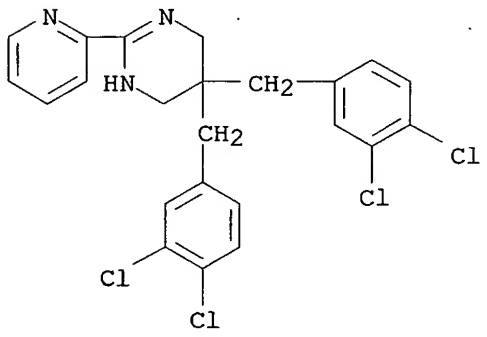

●2 HCl

RN 27337-19-5 CAPLUS
CN Pyrimidine, 5,5-bis(3,4-dichlorobenzyl)-1,4,5,6-tetrahydro-2-(3-pyridyl)-, dihydrochloride (8CI) (CA INDEX NAME)



●2 HCl

RN 27338-48-3 CAPLUS
CN Pyrimidine, 5,5-bis(3,4-dichlorobenzyl)-1,4,5,6-tetrahydro-2-(2-pyridyl)-, hydrochloride (8CI) (CA INDEX NAME)



●x HCl

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=> d his

     (FILE 'HOME' ENTERED AT 11:39:29 ON 24 SEP 2002)

     FILE 'REGISTRY' ENTERED AT 11:39:34 ON 24 SEP 2002
L1             STRUCTURE UPLOADED
L2          10 S L1 SSS SAM
L3             STRUCTURE UPLOADED
L4           0 S L3 SSS SAM
L5             STRUCTURE UPLOADED
L6           0 S L5 SSS SAM
L7          49 S L5 SSS FUL

     FILE 'CAPLUS' ENTERED AT 11:42:43 ON 24 SEP 2002
L8          20 S L7

     FILE 'CAOLD' ENTERED AT 11:43:39 ON 24 SEP 2002

=> s l7
L9           0 L7

=> log y
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                         ENTRY    SESSION
FULL ESTIMATED COST                                       0.38     230.96

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)          SINCE FILE      TOTAL
                                                         ENTRY    SESSION
CA SUBSCRIBER PRICE                                       0.00     -12.39

STN INTERNATIONAL LOGOFF AT 11:43:53 ON 24 SEP 2002
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